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## Author Correction: SIRT2-mediated deacetylation and deubiquitination of C/EBPB prevents ethanol-induced liver injury

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In the original publication of this article, we inadvertently misplaced an incorrect image for the 4-HNE (Pair-fed LoxP group) in Fig. 2a. The correct Fig. 2a is displayed as below. This correction does not affect the results or the conclusion of this work.

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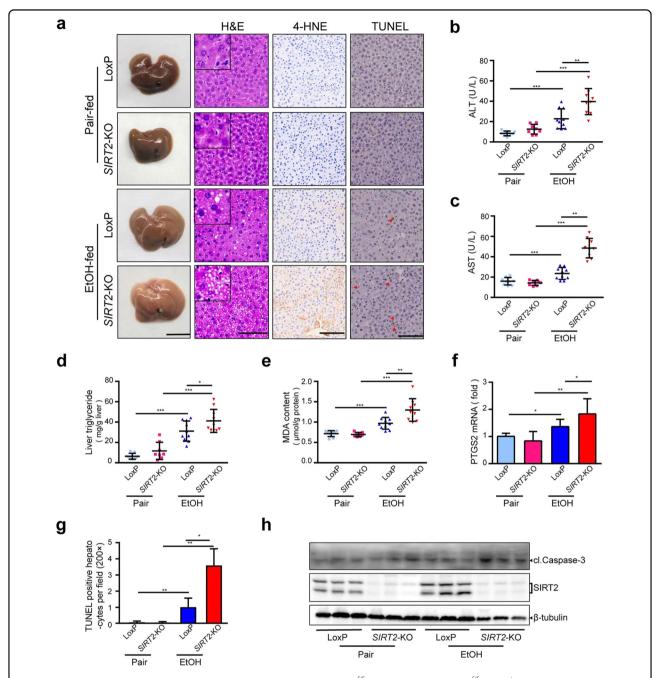


Fig. 2 Liver-specific SIRT2KO sensitizes mice to alcoholic liver injury.  $SIRT2^{ff}Alb$ - $Cre^-$  (LoxP) and  $SIRT2^{ff}Alb$ - $Cre^+$  (SIRT2-KO) male mice were treated with pair (Pair) and ethanol diet (EtOH) according to NIAAA model (n=8-10/group). **a-h** Liver injury, steatosis, lipid peroxidation, and cell apoptosis were assessed by images of the indicated livers (scale bar, 1 cm), mouse hepatic H&E staining (scale bar, 100  $\mu$ m), IHC detection of 4-HNE and TUNEL (scale bar, 100  $\mu$ m) (**a**), serum ALT (**b**) and AST (**c**), liver triglyceride (TG) (**d**), hepatic MDA content (**e**), and PTGS2 mRNA (**f**), quantitative analysis of TUNEL-positive hepatocytes (magnification,  $\times$ 200) (**g**), Western blot analysis of cl.Caspase-3 in murine liver tissues (**h**). Student's t-test was used for statistical evaluation. Data are shown as means  $\pm$  SD and are considered statistically significant at \*P < 0.05, \*\*P < 0.01 and \*\*\*P < 0.001.